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AMENDMENTS TO THE CLAIMS

This listing of claims will replace all prior versions and listings of claims in the application:

LISTING OF CLAIMS:

- 1. (canceled).
- 2. (currently amended): A compound according to claim 1, depicted by one of the general formulae (III) or (VIII):

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$$R_{1}$$
 R_{2}
 R_{3}
 R_{4}
 R_{2}
 R_{4}
 R_{4}
 R_{3}
 R_{5}
 R_{6}
 R_{6}
 R_{7}
 R_{8}
 R_{1}
 R_{2}
 R_{4}
 R_{3}
 R_{5}
 R_{6}
 R_{7}
 R_{8}
 R_{1}
 R_{2}
 R_{1}
 R_{2}
 R_{3}
 R_{4}
 R_{5}
 R_{6}
 R_{7}
 R_{8}
 R_{1}
 R_{2}
 R_{3}
 R_{4}
 R_{5}
 R_{6}
 R_{7}
 R_{8}

wherein

 R_1 , R_2 , R_3 , R_4 , R_5 , R_6 , R_7 and R_8 are independently hydrogen, hydroxy, OR_9 , $OC(O)R_9$, $OS(O)R_9$, alkyl, aryl, arylalkyl, thiol, alkylthio, bromo, chloro or fluoro, R_9 is alkyl, fluoroalkyl or arylalkyl,

 R_{13} — R_{14} and R_{15} are independently hydrogen, amino, cyano, thiol, nitro, or optionally substituted alkyl, haloalkyl, acyl, aryl, arylalkyl or alkylaryl, or the substituents R_{14} and R_{15} together with the nitrogen atom to which they are attached form an optionally substituted cyclic heteroalkyl or heteroaromatic structure,

 R_{46} — and R_{17} are independently hydrogen, amino, cyano, thiol, nitro or optionally substituted alkyl, haloalkyl, acyl, aryl, arylalkyl or alkylaryl, or the substituents R_{16} and R_{17} taken together

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with the carbon atom to which they are attached form an optionally substituted isoflavonoid ring system, and

the drawing "...." represents either a single bond or a double bond, which compounds include pharmaceutically acceptable salts thereof,

with the proviso that the following compounds

2,3-Dihydro-2,3-diphenyl-4H-1-benzopyran-4-one hydrazone

2'-Hydroxy-isoflavanone (2,4-dinitrophenyl)hydrazone

7-Methoxy-isoflavanone phenylhydrazone

5,7-Dimethoxy-isoflavanone (2,4-dinitrophenyl)hydrazone

Isoflavanone (2,4-dinitrophenyl)hydrazone

6-Hydroxy-isoflavanone (2,4-dinitrophenyl)hydrazone

7-Hydroxy-isoflavanone (2,4-dinitrophenyl)hydrazone

Isoflavanone semicarbazone

7-Methoxy-isoflavanone (2,4-dinitrophenyl)hydrazone

7-Hydroxy-4'-methoxy-isoflavanone (2,4-dinitrophenyl)hydrazone

5,7-Dimethoxy-isoflavanone (2,4-dinitrophenyl)hydrazone

6-Methoxy-isoflavanone (2,4-dinitrophenyl)hydrazone

4',5,7-trimethoxy-isoflavanone (2,4-dinitrophenyl)hydrazone

7-Methoxy-2-methyl-isoflavanone (2,4-dinitrophenyl)hydrazone

2-(Hydroxymethyl)-7-methoxy-isoflavanone (2,4-dinitrophenyl)hydrazone

and hydrochloride salts thereof are specifically excluded.

(currently amended): A compound according to claim <u>1</u>2, wherein R₁ is hydrogen,

 R_2 , R_3 , R_5 , R_6 and R_8 are independently hydrogen, hydroxy, OR_9 , $OC(O)R_9$, alkyl, aryl or arylalkyl,

R₄ and R₇ are independently hydroxy, OR₉ or OC(O)R₉,

 $\ensuremath{R_9}$ is methyl, ethyl, propyl, isopropyl or trifluoromethyl, and

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R₁₃,—R₁₄ and R₁₅ are independently hydrogen, methyl, ethyl, propyl, isopropyl, trifluoromethyl or optionally substituted phenyl, naphthyl or benzyl, or the substituents R₁₄ and R₁₅ together with the nitrogen atom to which they are attached form an optionally substituted cyclic heteroalkyl or heteroaromatic structure.

which compounds include pharmaceutically acceptable salts thereof,

 R_{16} —and R_{17} are independently hydrogen, methyl, ethyl, propyl, isopropyl, trifluoromethyl or optionally substituted phenyl, naphthyl or benzyl, or the substituents R_{16} and R_{17} taken together with the carbon atom to which they are attached form an optionally substituted isoflavonoid ring system, and

the drawing "-- " represents either a single bond or a double bond.

 (currently amended): A compound according to claim <u>2</u>3, wherein R₁ is hydrogen.

 $R_2,R_3,R_5,R_6 \ and \ R_8 \ are \ independently \ hydrogen, \ hydroxy, OR_9, OC(O)R_9 \ or \ methyl,$ $R_4 \ and \ R_7 \ are \ independently \ hydroxy, OR_9 \ or \ OC(O)R_9,$

Ro is methyl,

 R_{13} is hydrogen, methyl, ethyl, trifluoromethyl, phenyl, chlorophenyl, nitrophenyl, toluyl, naphthyl, benzyl, chlorobenzyl, nitrobenzyl or methylbenzyl,

R₁₄ is hydrogen and R₁₅ is hydrogen, methyl, ethyl, trifluoromethyl, phenyl, chlorophenyl, nitrophenyl, toluyl, naphthyl, benzyl, chlorobenzyl, nitrobenzyl or methylbenzyl, or the substituents R₁₄ and R₁₅ together with the nitrogen atom to which they are attached form an optionally substituted cyclic heteroalkyl or heteroaromatic structure,

 R_{16} —and R_{17} are independently hydrogen, methyl, ethyl, trifluoromethyl, phenyl, chlorophenyl, nitrophenyl, tohuyl, naphthyl, benzyl, chlorobenzyl, nitrobenzyl or methylbenzyl, or the substituents R_{16} and R_{17} taken together with the carbon atom to which they are attached form an optionally substituted isoflavonoid ring system, and

the drawing "--- " represents a single bond,

which compounds include pharmaceutically acceptable salts thereof.

(currently amended): A compound according to claim 34 selected from compounds (1) -

(10)(14):

4',7-Dihydroxyisoflavanone (phenyl)hydrazone (1)

4',7-Dihydroxyisoflavanone (4-nitrophenyl)hydrazone (2)

4'.7-Dihydroxvisoflavanone (4-methylphenyl)hydrazone (3)

4'.7-Dihydroxyisoflavanone (benzyl)hydrazone (4)

4'.7-Dihydroxyisoflavanone (4'.7-dihydroxyisoflavanone)hydrazone (5)

4',7-Dihydroxyisoflavanone (2-chlorophenyl)hydrazone (6)

4',7-Dihydroxyisoflavanone (3-chlorophenyl)hydrazone (7)

4'.7-Dihydroxvisoflavanone (4-chlorophenyl)hydrazone (8)

4',7-Dihydroxyisoflavanone (2-pyridyl)hydrazone (9)

4'.7-Dihydroxvisoflavanone (4-cyanophenyl)hydrazone (10)

4'.7 Dihydroxy 4 methylimino-isoflavan (11)

4'.7-Dihydroxyisoflavanone oxime (12)

4 Amino 3',4'-dimethoxy-7-hydroxy-8-methylisoflavan (13)

N-[3',4' dimethoxy-7-hydroxy-8-methyl-4-chromanyl) acetamide (14) which compounds include pharmaceutically acceptable salts thereof.

(currently amended): A process for the preparation of a compound of formula (III) or 6. (VIII)(1) as defined in claim 2 as claimed in claim 1 comprising the step of reacting the 4-keto group of a compound of the formula (X):

wherein

R2, R3, R4, R5, R6, R7, R8 and X are as defined in claim 1, and the drawing "---" represents either a single bond or a double bond,

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treatment, prophylaxis or amelioration of atherosclerosis.

with a hydrazine an-aminating agent.

7. (currently amended): A method for the treatment, prophylaxis or amelioration of a disease or disorder which method includes the step of administering a therapeutically effective amount of one or more compounds of formula (III) or (VIII)(+) as defined in claim 2 or a pharmaceutically acceptable salt or derivative thereof to a subject, with the proviso that the compounds and pharmaceutically acceptable salts of 3,4 Dihydro-3 phenyl-2H-1 benzopyran 4 amine N (3,4 Dihydro-3 phenyl-2H-1 benzopyran 4 yl) α phenyl benzeneacetamide, and N [3,4 Dihydro-3 phenyl-2H-1 benzopyran 4 yl) α phenyl benzeneacetamide, and N [3,4 Dihydro-3 phenyl-2H-1 benzopyran 4 yl) α phenyl benzeneacetamide, and N [3,4 Dihydro-3 phenyl-2H-1 benzopyran 4 yl) α phenyl-2H-1 benzeneacetamide, and N [3,4 Dihydro-3 phenyl-3 phe

3 (4 hydroxyphenyl) 2H-1 benzopyran 4 yll a phenyl benzeneacetamide are disclaimed for the

8. (currently amended): A method for the treatment, prevention or amelioration of diseases associated with aberrant cell survival, aberrant cell proliferation, abnormal cellular migration, abnormal angiogenesis, abnormal estrogen/androgen balance, dysfunctional or abnormal steroid genesis, degeneration including degenerative changes within blood vessel walls, inflammation, and immunological imbalance, which comprises administering to a subject one or more compounds of the formula (III) or (VIII)(t) as defined in claim 2 or a pharmaceutically acceptable salt or derivative thereof optionally in association with a carrier and/or excipient, with the provise that the compounds and pharmaceutically acceptable salts of

3,4 Dihydro 3 phenyl-2H-I benzopyran 4 amine N-(3,4 Dihydro 3 phenyl-2H-I benzopyran 4 yl) α phenyl-benzeneacetamide, and N-[3,4 Dihydro 3 (4 hydroxyphenyl) 2H-I benzopyran 4 yl] α phenyl-benzeneacetamide are disclaimed for the treatment, prophylaxic or amelioration of atherosclerosis.

- 9. (currently amended): A method of inducing apoptosis in cells expressing abnormal prosurvival phenotype which comprises contacting said cells with one or more compounds of the formula (III) or (VIII)(#) as defined in claim 2 or a pharmaceutically acceptable salt or derivative thereof optionally in association with a carrier or excipient.
- 10. (currently amended): A method for inhibiting migration of cells having an abnormal cellular

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migration phenotype which comprises contacting said cells with a compound of the formula (III) or (VIII)(F) as defined in claim 2 or a pharmaceutically acceptable salt or derivative thereof optionally in association with a carrier or excipient.

11. (currently amended): A method for inhibiting angiogenesis in tissue expressing aberrant angiogenic phenotype which comprises contacting said tissue with a compound of the formula (III) or (VIII)(f) as defined in claim 2 or a pharmaceutically acceptable salt or derivative thereof optionally in association with a carrier or excipients

with the proviso that the compounds and pharmaceutically acceptable salts of 3,4 Dihydro 3 phenyl-2H-1-benzopyran 4-amine
N (3,4 Dihydro 3-phenyl-2H-1-benzopyran 4-yl) α-phenyl-benzeneacetamide, and
N [3,4 Dihydro 3 (4 hydroxyphenyl)-2H-1-benzopyran 4-yl] α-phenyl-benzeneacetamide are disolaimed for the treatment, prophylaxis or amelioration of atherescelerosis

- 12. (currently amended): A method for the treatment, prevention or amelioration of cancer in a mammal which method comprises the step of bringing a compound of formula (III) or (VIII)(1) as defined in claim 2 or a pharmaceutically acceptable salt or derivative thereof into contact with cancerous tissue in a mammal that is suffering from a tumour, such that neoplastic development in said cancerous tissue is retarded or arrested.
- (canceled).
- (canceled).
- 15. (currently amended): An agent for the treatment, prophylaxis or amelioration of a disease or disorder, which agent comprises one or more compounds of formula (IIII) or (VIII)(4) as defined in claim 2 or a pharmaceutically acceptable salt or derivative thereof;

with the proviso that the compounds and pharmaceutically acceptable salts of 3,4

Dihydro 3 phenyl 2H-1 benzopyran 4 amine

N (3.4 Dihydro 3 phenyl 2H 1 benzopyran 4 yl) α phenyl benzeneacetamide, and

N [3.4 Dihydro 3 (4 hydroxyphenyl) 2H 1 benzopyran 4 yll α phenyl benzeneacetamide are

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disclaimed for the treatment, prophylaxis or amelioration of atheroselerosis.

16. (currently amended): A pharmaceutical composition which comprises one or more compounds of formula (III) or (VIII)(1) as defined in claim 2 or a pharmaceutically acceptable salt or derivative thereof in association with one or more pharmaceutical carriers, excipients, auxiliaries and/or diluents, with the proviso that the compounds and pharmaceutically acceptable salts of 3,4 Dihydro 3 phenyl 2H 1 benzopyran 4 amine N (3,4 Dihydro 3 phenyl 2H 1 benzopyran 4 yl) α-

phenyl benzeneacetamide, and N [3,4 Dihydro-3 (4 hydroxyphenyl) 2H 1 benzopyran 4 y1] α-

phenyl benzeneacetamide are disclaimed.

 (currently amended): A drink or food-stuff, which contains one or more compounds of formula (III) or (VIID(+) as defined in claim 2 or a pharmaceutically acceptable salt or derivative

thereof.

18. (currently amended): A compound of formula (III) or (VIII)(1) as defined in claim 2 or a

pharmaceutically acceptable salt thereof as herein described with reference to the Examples and/or

accompanying drawings.

19. (new) A compound selected from:

4',7-Dihydroxy-4-methylimino-isoflavan (11)

4',7-Dihydroxyisoflavanone oxime (12)

4-Amino-3',4'-dimethoxy-7-hydroxy-8-methylisoflavan (13)

N-[3',4'-dimethoxy-7-hydroxy-8-methyl-4-chromanyl)-acetamide (14)

which compounds include pharmaceutically acceptable salts thereof.

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